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Cost-effectiveness of self-management of blood pressure in hypertensive patients over 70 years with suboptimal control and established cardiovascular disease or additional cardiovascular risk diseases (TASMIN-SR)

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Abstract

Background: A previous economic analysis of self-management, that is, self-monitoring with self-titration of antihypertensive medication evaluated cost-effectiveness among patients with uncomplicated hypertension. This study considered cost-effectiveness of self-management in those with raised blood pressure plus diabetes, chronic kidney disease and/or previous cardiovascular disease.

Design and methods: A Markov model-based economic evaluation was undertaken to estimate the long-term cost-effectiveness of self-management of blood pressure in a cohort of 70-year-old 'high risk' patients, compared with usual care. The model used the results of the TASMIN-SR trial. A cost-utility analysis was undertaken from a UK health and social care perspective, taking into account lifetime costs of treatment, cardiovascular events and quality adjusted life years. A subgroup analysis ran the model separately for men and women. Deterministic sensitivity analyses examined the effect of different time horizons and reduced effectiveness of self-management.

Results: Base-case results indicated that self-management was cost-effective compared with usual care, resulting in more quality adjusted life years (0.21) and cost savings (–£830) per patient. There was a 99% chance of the intervention being cost-effective at a willingness to pay threshold of £20,000 per quality adjusted life year gained. Similar results were found for separate cohorts of men and women. The results were robust to sensitivity analyses, provided that the blood pressure lowering effect of self-management was maintained for more than a year.

Conclusion: Self-management of blood pressure in high-risk people with poorly controlled hypertension not only reduces blood pressure, compared with usual care, but also represents a cost-effective use of healthcare resources.

Keywords

Hypertension, self-management, cost-effectiveness, decision model, decision analysis

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Background

Hypertension is a leading risk factor for cardiovascular mortality and morbidity worldwide.^{1,2} Despite evidence of cost saving from antihypertensive treatment,³ and improvements in blood pressure (BP) monitoring, management and treatment,^{3,4} significant numbers of people remain inadequately controlled hence new models of care are required.⁵ Self-management of hypertension, in which an individual self-monitors their own BP and adjusts their own medication has been shown to lead to significantly lower BP in hypertension, including in those with higher cardiovascular risk.^{6,7}

The only economic analysis of self-management in the control of hypertension to date demonstrated that tele-monitoring with self-titration in uncomplicated hypertension was highly cost-effective, with incremental cost-effectiveness ratios below £5000 per quality adjusted life year (QALY) gained for men and women, when modelled over patient lifetime.⁸ However, subgroup analysis in the main trial suggested that the intervention might not be as effective in those with significant comorbidities, although patient numbers for this subgroup were small.⁷ Therefore, the TASMIN-SR trial was undertaken to determine the effect of self-monitoring with self-titration of antihypertensive medication on systolic BP among hypertensive patients with suboptimal control and pre-existing cardiovascular disease, diabetes mellitus (DM) and/or chronic kidney disease (CKD), compared with usual care. A model-based probabilistic cost-utility analysis was undertaken as part of this study to assess the long-term cost-effectiveness of the self-management intervention in a 'high risk' patient population, compared with usual care.

Methods

A Markov cohort model, built in TreeAge Pro (TreeAge Software Inc, Williamstown, MA, USA), was developed to estimate the long-term cost-effectiveness of self-management of BP compared with usual care, in patients with hypertension and a history of stroke, coronary heart disease (CHD), DM or CKD. The analysis used the results of the TASMIN-SR trial on BP, extrapolating these to long-term risk of cardiovascular endpoints (see below). Full details of the trial methods and results have been described in detail elsewhere.^{6,9} The model was run over a lifetime (30 year) time horizon using a six-month time cycle, with results presented from a UK National Health Service (NHS) and personal social services perspective.

Study population

The base case analysis considered a cohort of 70-year-old patients (39% women) with suboptimal hypertension,

BP $\geq 130/80$ mmHg at baseline, combined with a history of stroke, CHD, DM or CKD.⁶ Patients had at least one of four main underlying conditions (DM, stroke, CHD and CKD), to be eligible with 15 possible combinations of high risk conditions in total. Further details of the combined risk conditions are available in supplementary Table 2.

Interventions

Patients randomly assigned to usual care booked an appointment for a routine BP pressure check and medication review with the study general practitioner. Thereafter, usual care consisted of the participants seeing their general practitioner and or nurse for routine BP measurement and adjustment of medication at the discretion of the health professional. Patients randomly assigned to self-management were trained to self-monitor BP and to self-titrate their antihypertensive medication following a predetermined plan, in two or three sessions, each lasting around an hour. Following training, patients adjusted their antihypertensive medication based on their monthly self-monitored BP readings.⁹

Model structure

A patient entered the model in the high-risk health state and could move to another health state if they experienced one of three possible cardiovascular events (stroke, myocardial infarction (MI), unstable angina (UA)), or died from other causes (Figure 1). After a cardiovascular event, individuals could survive from that event or die within the first six months. Those who survived an event subsequently moved to a chronic health state for that condition until death, with no recurrences of cardiovascular events. For each chronic health state, an ongoing healthcare cost was applied every time cycle and quality of life was permanently reduced. Movement between health states was defined by transition probabilities, which represented the risk of experiencing an event within each six-month time cycle.

Model parameters

Patient level data from the TASMIN-SR trial were used to reflect the cardiovascular disease history of patients entering the Markov model. The probabilities of experiencing a stroke, MI or developing UA were obtained from published literature for hypertensive patients with each of the high-risk conditions^{10–14} (Table 1).^{15–17} When the model required probabilities that were not available in the literature (for a given age group, gender or combination of high risk conditions), missing values were estimated through extrapolation

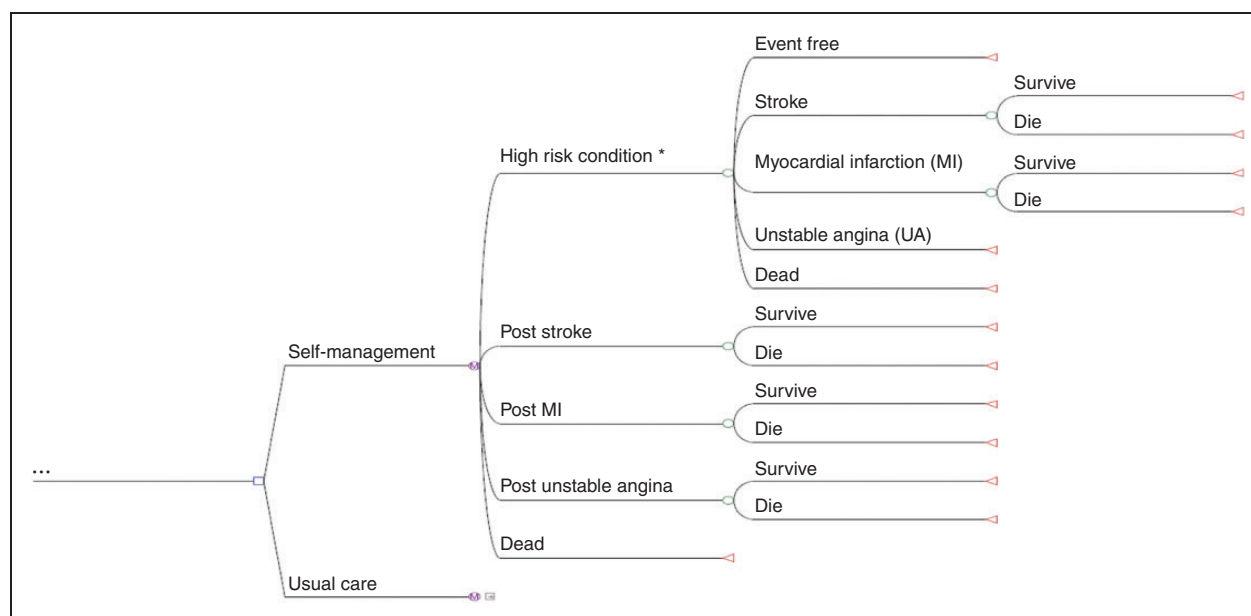


Figure 1. Markov model.

*In the Markov model patients can enter through any of fifteen combinations of high risk conditions.

Note: The Markov model is displayed for the self-management strategy only however, the model is identical for the usual care strategy.

(see supplementary document). For patients presenting with two or more high risk conditions, the probability of an event was calculated as the sum of the two individual risk probabilities (supplementary Tables 1 and 2).

Systolic BP reductions recorded in the trial at six months (11.4 mmHg and 5.5 mmHg for the intervention and control arms) and at 12 months (15.0 mmHg and 5.8 mmHg for the intervention and control arms) were extrapolated to age-related risk reductions for CHD (comprising both MI and UA) and stroke, using Law et al.¹⁸ (Table 1). Relative risks for CHD and stroke related to six and 12 month BP reductions are reported in Table 1. The model assumed that BP remained static for the first six-month cycle of the model, then reduced as per the six-month trial results for the second model cycle followed by the 12-month trial reductions thereafter with the between groups differences assumed constant in the base case. The probabilities of death from MI and stroke within a year of the event are reported in Table 1 and applied to the first year after an event (first two cycles in the model). Life tables were used to determine overall mortality, dependent on age and gender.¹⁹

Resource use and costs

Costs are reported in UK pounds at 2011/2012 prices. Resource use related to ongoing BP monitoring in primary care, self-management and prescription of antihypertensive agents was obtained from the TASMIN-SR trial at 12 months follow-up.

For self-management, equipment and training costs were annuitised at an annual rate of 3.5% and based on a lifetime of five years.²⁰ Replacement costs for the equipment and training were included at five yearly intervals over the lifetime of the model (see supplementary Table 3). Equipment used by individuals who died within any five-year interval was assumed to be discarded. Unit costs were applied to resource use and mean patient costs per six months were calculated for both randomly assigned groups, and applied to the initial high risk health state. Costs for acute and chronic cardiovascular event states were obtained from published studies^{14,21–23} (see Table 1).

Utility values

The primary outcome measure was QALYs. All utility scores used in the model are shown in Table 1. The utility values for the starting high risk health state were obtained from the TASMIN-SR trial, in which the overall mean EQ-5D score for hypertensive patients at baseline was used to estimate utilities. This was adjusted for age group using weights calculated from Ara and Brazier,²⁴ which allowed the overall reduction in quality of life with increasing age to be incorporated in the model. Acute events were assumed to happen approximately three months into a six-month cycle and individuals stayed in that acute state for three months before moving into a chronic state. Therefore, utilities for the acute state were applied mid-way through the six-month cycle and chronic health state

Table 1. Model parameters.

Parameter	Value	Source
Reduction in systolic BP at 12 months (mmHg)		TASMIN-SR trial ⁶
Self-management	15.0	
Usual care	5.8	
Reduction in systolic BP at 6 months (mmHg)		TASMIN-SR trial ⁶
Self-management	11.4	
Usual care	5.5	
Annual transition probabilities		
CVD events for patients with DM		NICE Diabetes guidelines, Appendix D1 ¹²
Stroke		
60–69 years old	0.0196	
70–79 years old	0.0262	
80–89 years old	0.0298	
MI		
60–69 years old	0.0089	
70–79 years old	0.0100	
80–89 years old	0.0111	
UA		
60–69 years old	0.0041	
70–79 years old	0.0047	
80–89 years old	0.0052	
CVD events for patients with CKD		Kerr et al. ¹¹
Stroke		
60–69 years old	0.0072	
70–79 years old	0.0147	
80–89 years old	0.0189	
MI		
60–69 years old	0.0051	
70–79 years old	0.0113	
80–89 years old	0.0171	
UA		
60–69 years old	0.0024	
70–79 years old	0.0054	
80–89 years old	0.0081	
CVD events for patients with a previous stroke		PROGRESS & NICE, lipid modification guidelines ^{10,14}
Stroke		
60–69 years old	0.0348	
70–79 years old	0.0589	
80–89 years old	0.0713	
MI		
60–69 years old	0.0139	
70–79 years old	0.0232	
80–89 years old	0.0232	
UA		
60–69 years old	0.0139	
70–79 years old	0.0232	
80–89 years old	0.0232	
CVD events for patients with CHD		NICE, lipid modification guidelines ¹⁴ and NICE hypertension guidelines ⁴

(continued)

Table 1. Continued

Parameter	Value	Source
Stroke		
60–69 years old	0.0359	
70–79 years old	0.0588	
80–89 years old	0.0713	
MI		
60–69 years old	0.0666	
70–79 years old	0.1112	
80–89 years old	0.1112	
UA		
60–69 years old	0.0528	
70–79 years old	0.0881	
80–89 years old	0.0881	
Age-related relative risks at 12 months (95% CI)		TASMIN-SR trial & Law et al. ^{6,18}
MI and UA – self-management		
60–69 years old	0.63 (0.60, 0.66)	
70–79 years old	0.68 (0.64, 0.71)	
80–89 years old	0.74 (0.70, 0.78)	
Stroke – self-management		
60–69 years old	0.53 (0.49, 0.57)	
70–79 years old	0.59 (0.55, 0.64)	
80–89 years old	0.74 (0.69, 0.79)	
MI and UA – usual care		
60–69 years old	0.83 (0.81, 0.84)	
70–79 years old	0.85 (0.84, 0.87)	
80–89 years old	0.89 (0.87, 0.90)	
Stroke – usual care		
60–69 years old	0.77 (0.75, 0.79)	
70–79 years old	0.81 (0.79, 0.83)	
80–89 years old	0.89 (0.86, 0.91)	
Age-related relative risks at 6 months (95% CI)		TASMIN-SR trial & Law et al. ^{6,18}
MI and UA – self-management		
60–69 years old	0.71 (0.68, 0.73)	
70–79 years old	0.75 (0.72, 0.77)	
80–89 years old	0.80 (0.76, 0.83)	
Stroke – self-management		
60–69 years old	0.62 (0.59, 0.66)	
70–79 years old	0.68 (0.64, 0.71)	
80–89 years old	0.80 (0.76, 0.84)	
MI and UA – usual care		
60–69 years old	0.83 (0.82, 0.85)	
70–79 years old	0.86 (0.85, 0.87)	
80–89 years old	0.89 (0.87, 0.91)	
Stroke – usual care		
60–69 years old	0.77 (0.75, 0.80)	
70–79 years old	0.81 (0.80, 0.84)	
80–89 years old	0.89 (0.87, 0.91)	
Probability of death for those who have experienced an event		
Fatal stroke	0.23	Bamford et al. ¹⁵

(continued)

Table 1. Continued

Parameter	Value	Source
Fatal MI		ONS, deaths registry 2011 & Kerr et al. ^{11,19}
65–74 years old	0.23	
75–84 years old	0.39	
85 and over	0.52	
Costs (UK £)		
Cost for the initial state ^a		TASMIN-SR trial, Curtis & BNF ^{6,16,17}
Self-management ^b	183	
Usual care	125	
Costs of acute disease one-off cost		
Stroke	11,020	Youman et al. ²³
MI	5487	Robinson et al. ²²
UA	3292	Assumed 60% of MI
Costs for long-term (chronic) disease per year		
Stroke	2721	Youman et al. ²³
MI	572	NICE, lipid modification guidelines ¹⁴
UA	572	NICE, lipid modification guidelines ¹⁴
Utilities		
Utilities for initial health state		
Self-management and usual care		TASMIN-SR trial ⁶
65–74 years old	0.81	
75–84 years old	0.74	
85 and over	0.71	
Utilities for acute events		NICE, lipid modification guidelines ¹⁴
UA	0.77	
MI	0.76	
Stroke	0.63	
Utilities for long-term (chronic) disease		NICE, lipid modification guidelines ¹⁴
UA	0.88	
MI	0.88	
Stroke	0.63	
Dead	0.00	By definition

^aIncluded annual costs of drugs per patient, average general practitioner and practice nurse cost of consultation(s) and the costs of the intervention. The cost difference between self-monitoring and usual care was driven by the cost of the intervention ^bFor greater detail see supplementary document. CVD: cardiovascular disease; CKD: chronic kidney disease; DM: diabetes mellitus; MI: myocardial infarction; UA: unstable angina.

utilities were applied at the start of the subsequent cycle (Table 1). Health state utilities for cardiovascular events were applied multiplicatively to the age-related high risk health state utility values.

Analysis

A cost–utility analysis was undertaken from a UK NHS and personal social services perspective. For the base-case analysis, 15 separate cost-effectiveness analyses were run, one for each combination of high risk conditions assessed in the model. The final cost-effectiveness results correspond to the trial population-weighted average of costs and QALYs and are reported in terms of the

incremental cost per QALY gained.²⁵ Analyses were also separately run for men and women. Costs and outcomes were discounted at an annual rate of 3.5%.²⁶

Uncertainty in the model results was assessed using sensitivity analyses. Deterministic sensitivity analysis was undertaken around key parameters and assumptions. The time horizon for the model was varied from 30 years (lifetime) to between one year and 20 years, to determine whether the intervention was cost-effective in the shorter term. The assumption regarding the long-term effectiveness of the intervention was tested by assessing the impact of limiting the additional effect on BP lowering to years of self-management.^{1,2,5,10} Additional sensitivity analyses altered long-term

Table 2. Results of cost-effectiveness analysis.

	Costs	QALYs	Incremental cost	Incremental QALYs	ICER
Total population					
Usual care	8187	6.0326			
Self-management	7357	6.2466	−830	0.2139	Dominant
Women					
Usual care	7338	6.2467			
Self-management	6579	6.4456	−759	0.1988	Dominant
Men					
Usual care	8654	5.9035			
Self-management	7791	6.1257	−864	0.2221	Dominant

QALYs: quality adjusted life years; ICER: Incremental Cost-Effectiveness Ratio.

cardiovascular event costs by 30% (up and down). Finally, all analyses were re-run using the unadjusted trial data, which showed marginally smaller reductions in BP (11.4 mmHg and 5.8 mmHg for the intervention and control arms at six months and 14.9 mmHg and 6.0 mmHg, respectively, at 12 months). When possible, data were entered into the model as distributions in order that a probabilistic sensitivity analysis could be undertaken to incorporate parameter uncertainty. Gamma distributions were fitted to all costs obtained from the TASMIN-SR trial and beta distributions were applied to the utility values. The parameters used for these distributions are shown in Table 1. The probabilistic sensitivity analysis was run with 10,000 second-order Monte Carlo simulations and cost-effectiveness planes and cost-effectiveness acceptability curves constructed to estimate the probability of self-management being cost-effective at different willingness-to-pay thresholds.²⁰

Results

In the base-case analysis, self-management of BP was dominant compared to usual care, being cheaper and more effective (Table 2). Self-management was associated with mean cost savings of £830 per patient for the total population (self-management £7357 vs. usual care £8187) and a gain of 0.21 QALYs (6.25 vs. 6.03, respectively). This dominance was demonstrated for both men and women (Table 2). In the cost-effectiveness plane (Figure 2), all results are in the north-east and south-east quadrants indicating that self-management is always more effective but with greater uncertainty around the difference in costs. The cost-effectiveness acceptability curve shows that the probability of self-management of BP being cost-effective compared with usual care was at least 99% if decision-makers were willing to pay £20,000 per QALY gained. At a lower threshold of £10,000 per QALY, the probability of the intervention being cost-effective compared with usual care was still high at 97% (Figure 2).

A sensitivity analysis of time horizon demonstrated that self-management is dominant if the horizon is two years or more (Table 3). Similarly, if the impact of self-management on BP is time limited, the cost-effectiveness is reduced – but the intervention is still cost-effective provided that the effect is sustained for one year (first two cycles) (Table 4). Other sensitivity analyses (costs and reduced impact on BP) did not change the overall results (see supplementary Tables 4–6).

Discussion

This is the first study to present results of the cost-effectiveness of self-management of BP compared with usual care in a high risk population with suboptimally managed hypertension and significant cardiovascular comorbidity. The base-case analysis suggests that self-management of BP is cost-effective and is likely to be dominant (i.e. it is less costly and produces more QALYs) compared to usual care.

The main driver of this result is the estimated decline in the risk of cardiovascular events associated with the observed additional BP lowering achieved with self-management, and this explanation also holds for the greater benefit seen for men. This result was robust to sensitivity analysis unless the time horizon was reduced below two years or the observed BP lowering effect of self-management did not continue beyond a year.

Relationship with other literature

Previous economic studies have evaluated the cost-effectiveness of self-monitoring rather than self-management (self-monitoring plus self-titration of antihypertensive medication) and only one previous economic analysis of self-management has been undertaken (TASMINH2),⁸ which found self-management to be cost-effective (£1624 and £4923 per QALY gained for men and women, respectively).⁸ In this analysis, we

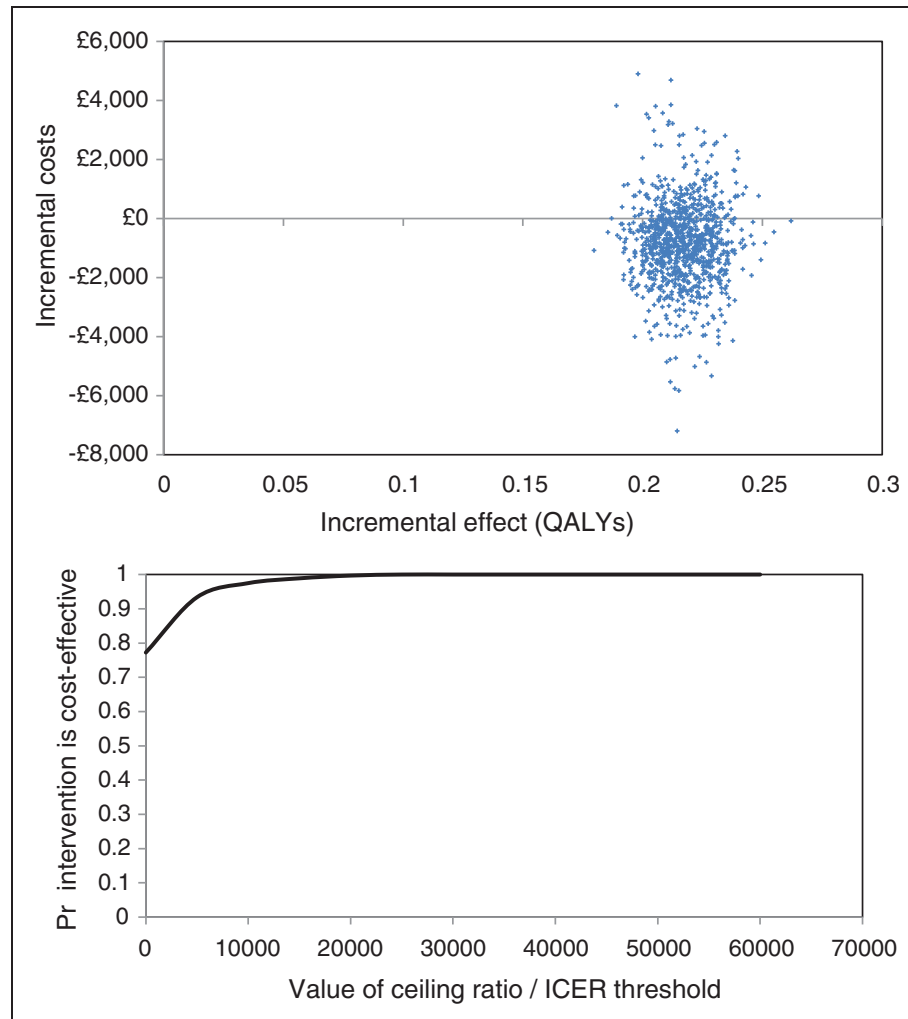


Figure 2. Base-case results.

Incremental cost-effectiveness plane: self-management against usual care. CEAC: cost-effectiveness acceptability curve. CEAC for self-monitoring of hypertension

found self-management to be even more cost-effective, reflecting the higher number of cardiovascular events predicted to have been prevented in the higher risk population, and the slightly greater reductions in BP that were observed in the TASMIN-SR trial.

Strengths and limitations

This study used cost and outcome data of trial participants,⁶ who may differ from similar patients not taking part in the trial for instance being more adherent and healthier.²⁷ The strongly positive results, however, suggest that such an intervention would be cost-effective even in a less compliant population. The costs of long-term and acute care were taken from estimates in the literature and a number of assumptions were made about the annual probabilities of cardiovascular events by risk conditions based on best published

information. A key assumption was that of the prolonged effectiveness of the intervention. In both TASMINH2 and TASMIN-SR, the difference in BP reduction between trial arms continued to diverge between six and 12 months, suggesting that the effect may be maintained over time. Indeed, an 18 month post-trial follow-up of the HSM self-management trial found that BP continued to diverge over time, suggesting our assumption of maintenance of effect may even be conservative.²⁸ The sensitivity analyses showed that even if BP differences lasted only one further year and then returned to the effectiveness of usual care, self-management is still likely to be cost-effective. For simplicity, the model did not include subsequent cardiovascular events. Given that the main driver of costs was events and the main driver of events was BP, it would be expected that a model including secondary and subsequent events would show self-management to be even

Table 3. Sensitivity analyses: cost-effectiveness by time horizon.

	Costs	QALYs	Incremental cost	Incremental QALYs	ICER
20 years					
Usual care	7709	5.8830			
Self-management	6919	6.0975	-789	0.2145	Dominant
10 years					
Usual care	5242	4.7756			
Self-management	4675	4.9252	-567	0.1496	Dominant
5 years					
Usual care	2882	3.1178			
Self-management	2554	3.1742	-328	0.0564	Dominant
3 years					
Usual care	1690	2.0859			
Self-management	1535	2.1044	-155	0.0186	Dominant
2 years					
Usual care	1116	1.4651			
Self-management	1056	1.4718	-59	0.0067	Dominant
1 year					
Usual care	603	0.7729			
Self-management	625	0.7736	22	0.0006	34,791

QALYs: quality adjusted life years; ICER: Incremental Cost-Effectiveness Ratio.

Table 4. Sensitivity analyses: cost-effectiveness by reducing the additional effect of self-management to blood pressure lowering at four different time points.

Time horizon	Costs	QALYs	Incremental cost	Incremental QALYs	ICER
10 years					
Usual care	8187	6.0326			
Self-management	7530	6.2242	-657	0.1916	Dominant
5 years					
Usual care	8187	6.0326			
Self-management	7876	6.1623	-311	0.1297	Dominant
2 years					
Usual care	8187	6.0326			
Self-management	8259	6.0757	71	0.0430	1660
1 year					
Usual care	8187	6.0326			
Self-management	8382	6.0454	195	0.0127	15,341

QALYs: quality adjusted life years; ICER: Incremental Cost-Effectiveness Ratio.

more cost-effective than usual care. The model considers patients with comorbidities and additional risk factors (e.g. age, gender). Arguably, a more complex model such as individual patient level simulation could be more appropriate in this situation, as this type of model can incorporate patient history more efficiently, overcoming the limitations of Markov

models.²⁹ Finally, an assumption has been made regarding the differential effect of BP lowering between the intervention and control groups. Systematic reviews suggest that lowering BP below 140/90 mmHg is as effective as lowering BP to 140/90 mmHg,¹⁸ but it is fair to say that the evidence of benefit is stronger in stroke and DM than in CHD or CKD.^{10,30-32}

Clinical implications

These results suggest that the benefits of BP reduction seen in the trial can be achieved in a highly cost-effective manner. The up-front costs of implementation of self-management of hypertension in high risk groups are relatively modest (£14.6 equipment and £20.0 training), and are soon repaid by future maintenance of quality of life and reductions in costs from reduced cardiovascular events. The very high likelihood of cost-effectiveness from both this and the previous analyses suggest that self-management is a strong candidate for implementation.

Conclusions

The results of this model-based economic evaluation suggest that self-management of hypertension in high risk patients is a cost-effective strategy in the short and long term, resulting in QALY gains and cost savings. Self-management of BP in high risk patients represents an important new addition to the management of hypertension in primary care.

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Declaration of conflicting interests

The author(s) declared the following potential conflicts of interest with respect to the research, authorship, and/or publication of this article: RJM has received research equipment from Omron and Lloyds Pharmacies. BW has worked in academic collaboration with Healthstats, Singapore, in developing novel blood pressure monitoring approaches. FDRH has received limited research support in terms of blood pressure devices from Microlife and BP TRU. No other disclosures were reported.

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References

1. Lewington S, Clarke R, Qizilbash N, et al. Age-specific relevance of usual blood pressure to vascular mortality: a meta-analysis of individual data for one million adults in 61 prospective studies. *Lancet* 2002; 360: 1903–1913.
2. Lozano R, Naghavi M, Foreman K, et al. Global and regional mortality from 235 causes of death for 20 age groups in 1990 and 2010: a systematic analysis for the Global Burden of Disease Study 2010. *Lancet* 2012; 380: 2095–2128.
3. National Institute for Health and Clinical Excellence. *Hypertension: The clinical management of primary hypertension in adults. NICE clinical guideline 127*. London: NICE, 2011.
4. National Institute for Health and Clinical Excellence. *Hypertension: Management of hypertension in adults in primary care: Partial update. NICE guideline CG34*. London: NICE, 2006.
5. Falaschetti E, Mindell J, Knott C, et al. Hypertension management in England: a serial cross-sectional study from 1994 to 2011. *Lancet* 2014; 383: 1912–1919.
6. McManus RJ, Mant J and Haque MS. Effect of self-monitoring and medication self-titration on systolic blood pressure in hypertensive patients at high risk of cardiovascular disease: The TASMIN-SR randomized clinical trial. *JAMA* 2014; 312: 799–808.
7. McManus RJ, Mant J, Bray EP, et al. Telemonitoring and self-management in the control of hypertension (TASMINH2): a randomised controlled trial. *Lancet* 2010; 376: 163–172.
8. Kaambwa B, Bryan S, Jowett S, et al. Telemonitoring and self-management in the control of hypertension (TASMINH2): a cost-effectiveness analysis. *Eur J Prev Cardiol* 2014; 21: 1517–1530.
9. O'Brien C, Bray EP, Bryan S, et al. Targets and self-management for the control of blood pressure in stroke and at risk groups (TASMIN-SR): protocol for a randomised controlled trial. *BMC Cardiovasc Disorders* 2013; 13: 21.
10. PROGRESS Collaborative Group. Randomised trial of a perindopril-based blood-pressure-lowering regimen among 6105 individuals with previous stroke or transient ischaemic attack. *Lancet* 2001; 358: 1033–1041.
11. Kerr M, Bray B, Medcalf J, et al. Estimating the financial cost of chronic kidney disease to the NHS in England. *Nephrol Dial Transplant* 2012; 27(suppl 3): iii73–iii80.
12. National Institute for Health and Clinical Excellence. *National guidelines for the management of blood glucose levels in people with type 2 diabetes. NICE guideline CG87*. London: NICE, 2002.
13. National Institute for Health and Clinical Excellence. *Statins for the prevention of cardiovascular events. NICE guideline TA94*. London: NICE, 2006.
14. National Institute for Health and Clinical Excellence. *Lipid modification: Cardiovascular risk assessment and*

- the modification of blood lipids for the primary and secondary prevention of cardiovascular disease. NICE guideline CG67.* London: NICE, 2008.
15. Bamford J, Sandercock P, Dennis M, et al. A prospective study of acute cerebrovascular disease in the community: the Oxfordshire Community Stroke Project – 1981–86. 2. Incidence, case fatality rates and overall outcome at one year of cerebral infarction, primary intracerebral and subarachnoid haemorrhage. *J Neurol, Neurosurg Psychiatry* 1990; 53: 16–22.
 16. Joint Formulary Committee. *British National Formulary.* London: BMJ Publishing Group and RPS Publishing, 2012.
 17. Curtis L. *Unit Costs of Health and Social Care.* Kent: Personal Social Services Research Unit, University of Kent, 2012.
 18. Law MR, Morris JK and Wald NJ. Use of blood pressure lowering drugs in the prevention of cardiovascular disease: meta-analysis of 147 randomised trials in the context of expectations from prospective epidemiological studies. *BMJ* 2009; 338: b1665.
 19. Office for National Statistics. *Interim Life Tables for England, Newport,* Office for National Statistics, 2011.
 20. Gray AM, Clarke PM, Wolstenholme JL, et al. *Applied Methods of Cost-Effectiveness Analysis in Health Care.* Oxford: Oxford University Press, 2011.
 21. Department of Health. *NHS Reference Costs Schedule 2010–11.* London: Department of Health, 2013.
 22. Robinson M, Palmer S, Sculpher M, et al. Cost-effectiveness of alternative strategies for the initial medical management of non-ST elevation acute coronary syndrome: systematic review and decision-analytical modelling. *Health Technol Assess (Winchester, England)* 2005; 9iii–iv, ix–xi, 1–158.
 23. Youman P, Wilson K, Harraf F, et al. The economic burden of stroke in the United Kingdom. *Pharmacoeconomics* 2003; 21: 43–50.
 24. Ara R and Brazier JE. Using health state utility values from the general population to approximate baselines in decision analytic models when condition-specific data are not available. *Value in Health: The journal of the International Society for Pharmacoeconomics and Outcomes Research* 2011; 14: 539–545.
 25. Drummond MF, Sculpher MJ, Torrance GW, et al. *Methods for the economic evaluation of health care programmes,* 3rd ed. Oxford: Oxford University Press, 2005.
 26. National Institute for Health and Clinical Excellence. *Guide to the methods of technology appraisal. Process and methods guides.* London: NICE, 2013.
 27. Nallamothu BK, Hayward RA and Bates ER. Beyond the randomized clinical trial: the role of effectiveness studies in evaluating cardiovascular therapies. *Circulation* 2008; 118: 1294–1303.
 28. Maciejewski ML, Bosworth HB, Olsen MK, et al. Do the benefits of participation in a hypertension self-management trial persist after patients resume usual care? *Circ Cardiovasc Qual Outcomes* 2014; 7: 269–275.
 29. Barton P, Bryan S and Robinson S. Modelling in the economic evaluation of health care: selecting the appropriate approach. *J Health Serv Res Policy* 2004; 9: 110–118.
 30. Bangalore S, Kumar S, Volodarskiy A, et al. Blood pressure targets in patients with coronary artery disease: observations from traditional and Bayesian random effects meta-analysis of randomised trials. *Heart* 2013; 99: 601–613.
 31. Reboli G, Gentile G, Angeli F, et al. Effects of intensive blood pressure reduction on myocardial infarction and stroke in diabetes: a meta-analysis in 73,913 patients. *J Hypertens* 2011; 29: 1253–1269.
 32. Upadhyay A, Earley A, Haynes SM, et al. Systematic review: blood pressure target in chronic kidney disease and proteinuria as an effect modifier. *Ann Intern Med* 2011; 154: 541–548.